Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1479.

**SUPPLEMENTARY INFORMATION:** In compliance with section 3507 of the PRA (44 U.S.C. 3507), FDA has submitted the following proposed collection of information to OMB for review and clearance:

### Notice of Availability of Sample Electronic Product—21 CFR Parts 1020, 1030, 1040, and 1050 and FDA Form 2767 (OMB Control No. 0910–0048—Reinstatement)

Under sections 532 through 542 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360ii through 360ss), FDA regulates electronic products that emit radiation. Section 532 of the act directs the Secretary of the Department of Health and Human Services (the Secretary) to establish and carry out an electronic product radiation control program designed to protect the public health and safety from electronic radiation, and authorizes the Secretary to procure (by negotiation or otherwise) electronic products for research and testing purposes and to sell or otherwise dispose of such products.

FDA's Center for Devices and Radiological Health (CDRH) conducts laboratory compliance testing of products covered by regulations for product standards in parts 1020, 1030, 1040, and 1050 (21 CFR parts 1020, 1030, 1040, and 1050). The ''Notice of Availability of Sample Electronic Product'' (Form FDA 2767) is used to inform CDRH of the location of sample products that are being requested for testing to confirm that the products comply with performance standards. Form FDA 2767 is a summary form which reports information as required by parts 1020, 1030, 1040, and 1050.

FDA also uses this information to locate and select sample products to ensure conformance with regulations. In the event this information were not collected by CDRH, each manufacturer would have to respond in letter format with all the data now being recorded on Form FDA 2767, which would require more time and expense. Testing an appropriate percentage of these products to protect the public would also be hindered by the slower process.

The respondents to this collection of information are manufacturers of electronic products.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN

| 21 CFR Part and Form Number | No. of Respondents | Annual Frequency per Response | Total Annual Responses | Hours per Response | Total Hours |
|---|---|---|---|---|---|
| 1020, 1030, 1040, 1050, and Form FDA 2767 | 145 | 11.03 | 1,600 | 0.09 | 144 |
| Totals | | | | | 144 |

There are no capital costs or operating and maintenance costs associated with this collection of information.

FDA's estimates are based on actual data collected from industry over the past 3 years, where there has been an average of 1,600 annual responses to FDA from 145 respondents each year.

Dated: September 9, 1997.

**William K. Hubbard,**

*Associate Commissioner for Policy Coordination.*

[FR Doc. 97–24581 Filed 9–16–97; 8:45 am]

**BILLING CODE 4160–01–F**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Care Financing Administration

**[HCFA–R–52]**

### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Health Care Financing Administration, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimate burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

*Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Conditions for Coverage of Supplier of End Stage Renal Disease (ESRD) Services and Supporting Regulations Contained in 42 CFR 405.2100–2171; *Document No.:* HCFA–R–52 (OMB#0938–0386); *Use:* These conditions of coverage are needed to ensure proper distribution and effective utilization of ESRD treatment sources. In addition, the conditions maintain and improve the efficient delivery of care by physicians and dialysis facilities; *Frequency:* Annually; *Affected Public:* Business or other for-profit, and Federal Government; *Number of Respondents:* 2,976; *Total Annual Responses:* 2,976, *Total Annual Hours:* 100,937.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, E-mail your request, including your address, phone number, OMB number, and HCFA document identifier to Paperwork@hcfa.gov, or call the Reports Clearance Office on (410) 786–1326. Written comment and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Information Services, Information Technology Investment Management Group, Division of HCFA Enterprise Standards, Attention: Louis Blank, Room C2–26–17, 7500 Security Boulevard, Baltimore, Maryland, 21244–1850.

Dated: September 10, 1997.

**John P. Burke III,**

*HCFA Reports Clearance Officer, HCFA Office of Information Services, Information Technology Investment Management Group, Division of HCFA Enterprise Standards.*

[FR Doc. 97–24652 Filed 9–16–97; 8:45 am]

**BILLING CODE 4120–03–M**